

# CLASSIFICATION: HEAVY CONSTANT CHAINS (cont'd)

- 97) MOPC173: MOUSE IGG2A
- 98) CBPC101: MOUSE IGG2A
- 99) IGA'CL: MOUSE IGA
- 100) IGA'CL: MOUSE IGA
- 101) MOPC47A: MOUSE IGA
- 102) MOPC315: MOUSE IGA
- 103) MOPC311: MOUSE IGA
- 104) IGE'CL: MOUSE IGE
- 105) IGE'CL: MOUSE IGE
- 106) IGE a'CL: MOUSE IGE
- 107) IGE b'CL: MOUSE IGE
- 108) IGE MEMB'CL: MOUSE IGE MEMBRANE BOUND
- 109) IR-731: RAT IGD
- 110) RAT IGG2c'CL: RAT IGG2c
- 111) RAT IGG2a'CL: RAT IGG2a
- 112) RAT IGG1'CL: RAT IGG1
- 113) RAT IGG2b'CL: RAT IGG2b
- 114) IR2'CL: RAT IGE
- 115) PK3'CL: RABBIT MU CHAIN SECRETED (ALLOTYPE A2)
- 116) RABBIT IGG: RABBIT IGG
- 117) PCAN001-12,14'CL: RABBIT IGG
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## GENERAL NOTES: HEAVY CONSTANT CHAINS

# CH1, HINGE, CH2, CH3 AND CH4 DOMAINS ARE TABULATED TO CONFORM TO THEIR CODING NUCLEOTIDE SEQUENCES RELATIVE TO INTERVENING SEQUENCES ESTABLISHED BY SAKANO, H., ROGER, J.H., HUPPI, K., BRACK, C., TRAUNECKER, A., MAKI, R., WALL, R. & TONEGAWA, S. (1979) NATURE, 277, 627-633. A MEMBRANE DOMAIN IS LISTED SEPARATELY TO INCLUDE PART OF THE C-TERMINAL PORTION OF MEMBRANE BOUND IMMUNOGLOBULINS. IF IT IS DESIRED TO ALIGN THE LIGHT CHAIN WITH THE HEAVY CHAIN DOMAINS FOR HOMOLOGY, THE SEQUENTIAL NUMBERING IN THE FIRST COLUMN SHOULD BE USED: RESIDUES 108 TO 215 FOR CL; 114 TO 223 IN CH1 PLUS THE FIRST PART OF HINGE (224 TO 241), THE END OF HINGE (242 AND 243) AND THE FIRST TWO RESIDUES OF CH2 (244 AND 245); 246 THROUGH 361 OF CH2; 362 THROUGH 496 OF CH3; AND 497 THROUGH 628 OF CH4. GAPS IN THE SEQUENTIAL NUMBERING ARE USED FOR ALIGNMENT.

# DISULFIDE BONDS ARE LOCATED AT THE FOLLOWING POSITIONS IF CYS IS PRESENT:

INTRACHAIN:	142-208, 274-340, 249-312, 390-456, 524-587.
HL-INTERCHAIN:	127 OR 128, 198 OR 225, 230, 235.
HH-INTERCHAIN:	232, 233, 237, 238, 239, 240, 241A, G, M, P, V, BB, EE, KK, QQ, 242, 248, 261, 314.
INTERSUBUNIT:	328, 444.
TO J-CHAIN:	495, 627.

IDENTIFICATIONS OF SOME OF THESE DISULFIDE BONDS ARE NOT ABSOLUTELY CERTAIN.

# THERE WOULD APPEAR TO BE POLYMORPHISM AMONG GAL, OU, SCO AND BOT/CO MU-CHAINS AS FOLLOWS:

GAL	SER-334, VAL-358
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## SPECIFIC NOTES: HEAVY CONSTANT CHAINS

- 1) HUMAN IGM'CL: THE AMINO ACID SEQUENCE WAS OBTAINED BY TRANSLATING A CLONE OF HUMAN FETAL LIVER DNA.
- 5) BOT: FROM A CASE OF IGM HEAVY CHAIN DISEASE. THE AMINO ACID RESIDUES AT POSITIONS 451 TO 476, 519 TO 544 AND 508 TO 518 ARE IDENTICAL TO THAT OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN DNA CONTAINING THE IGM GENE (TAKAHASHI, N., NAKAI, S. & HONJO, T. (1980) NUCLEIC ACIDS RES. 8, 5983-5991).
- 7) GLI: IT HAS NO VARIABLE REGION, AND WAS FROM A CASE OF HEAVY CHAIN DISEASE.
- 9) WAB: THE SEQUENCES OF WAB AND NIG-65 ARE IDENTICAL FOR POSITIONS 241 PP TO 485 WHERE BOTH SEQUENCES ARE DETERMINED. FOR WAB, THERE ARE FOUR OR FIVE N-ACETYL-D-GALACTOSAMINE OLIGOSACCHARIDES ATTACHED TO SER AT POSITION 234, AND THR AT POSITIONS 241 J, 241 K, AND 241 O AND/OR 241 P. FOR NIG-65, THERE ARE THREE N-ACETYL-D-GLUCOSAMINE OLIGOSACCHARIDES ATTACHED TO ASN AT POSITIONS 314, 414 AND 468.
- 10) NIG-65: THE SEQUENCES OF WAB AND NIG-65 ARE IDENTICAL FOR POSITIONS 241 PP TO 485 WHERE BOTH SEQUENCES ARE DETERMINED. FOR WAB, THERE ARE FOUR OR FIVE N-ACETYL-D-GALACTOSAMINE OLIGOSACCHARIDES ATTACHED TO SER AT POSITION 234, AND THR AT POSITIONS 241 J, 241 K, AND 241 O AND/OR 241 P. FOR NIG-65, THERE ARE THREE N-ACETYL-D-GLUCOSAMINE OLIGOSACCHARIDES ATTACHED TO ASN AT POSITIONS 314, 414 AND 468.
- 11) ERI: THERE ARE THREE AMINO ACID RESIDUES BETWEEN POSITION 140 AND POSITION 142; THEY ARE ALA, VAL AND ALA. THE AUTHOR HAS RECENTLY INDICATED THAT THERE IS ONLY ONE AMINO ACID RESIDUE THERE AND IT IS ALA.
- 14) HUMAN IGG3'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF THREE CLONES OF HUMAN GENOMIC DNA. THERE ARE FOUR EXONS: POSITIONS 216 TO 241B, 241C TO 241O, 241R TO 241FF, AND 241GG TO 243.
- 15) OMH'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN CELL LINE CDNA.
- 16) HER: THE SEQUENCES OF HER, FRO AND JON ARE IDENTICAL.
- 17) FRO: THE SEQUENCES OF HER, FRO AND JON ARE IDENTICAL.
- 18) JON: THE SEQUENCES OF HER, FRO AND JON ARE IDENTICAL.
- 19) WIS: FROM A CASE OF HEAVY CHAIN DISEASE.
- 20) SPA: FROM A CASE OF HEAVY CHAIN DISEASE.
- 21) ZUC: FROM A CASE OF HEAVY CHAIN DISEASE.
- 22) ZUC: OBTAINED FROM THE SAME PATIENT AS ZUC, AND EXISTED IN A MONOMER FORM.
- 23) KUP: THE SEQUENCES OF KUP AND BRU ARE IDENTICAL.
- 24) BRU: THE SEQUENCES OF KUP AND BRU ARE IDENTICAL.
- 26) CRA: IT HAS NO VARIABLE REGION AND NO CHI REGION, AND WAS FROM A CASE OF HEAVY CHAIN DISEASE. AMINO ACID RESIDUES FOUND AT POSITION 224 ARE THR AND SER.
- 29) NIE: IN EARLIER EDITIONS, THE AMINO ACID RESIDUES AT POSITIONS 238, 285, 300 AND 331 LISTED BY US WERE INCORRECT.
- 30) CRA: FROM A CASE OF HEAVY CHAIN DISEASE.
- 31) VAO: FROM A CASE OF HEAVY CHAIN DISEASE.
- 32) LEB: FROM A CASE OF HEAVY CHAIN DISEASE.
- 33) EST: IT HAS NO VARIABLE REGION AND NO CHI REGION, AND WAS FROM A CASE OF HEAVY CHAIN DISEASE.
- 34) YOK: THE GLY AT POSITION 462 IN IGG1 CHAIN IS CONSIDERED TO BE ASSOCIATED WITH THE GM(2) ALLOTYPE (COOK, E. & STEINBERG, A.G. (1979) MOL. IMMUNOL. 16, 555-558).
- 36) HUMAN IGG1'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A HUMAN FETAL LIVER DNA.
- 41) BDR: THE AMINO ACID RESIDUES AT POSITIONS 262, 263, 269, 304, 310, 333, 430, 431, 447, 461 AND 490 HAVE RECENTLY BEEN REVISED BY THE AUTHOR. VITTHIAN, F., LUY, S., LOM, J. K. (1979) J. BIOL. CHEM. 254, 2865-2874. FROM GLN, ASP, GLU, GLN, ASP, ASN, GLN, GLY, ASP, GLU AND GLU TO GLX, ASX, GLX, GLX, ASX, GLX, GLN, ASX, GLX AND GLN RESPECTIVELY.
- 43) CAR: THE SEQUENCE WAS OBTAINED AFTER DIGESTION WITH PROTEASES. PROTEASES FROM S. SANGUIS AND S. PNEUMONIAE CLEAVED BETWEEN POSITIONS 241C AND 241D, AND THAT FROM H. INFLUENZAE BETWEEN POSITIONS 241G AND 241H.
- 47) HUMAN IGG'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN FETAL LIVER DNA. THIS GENE IS NOT EXPRESSED DUE TO THE LACK OF A SWITCH REGION, AND IS THUS DESIGNATED AS PSEUDO-GAMMA IGHGP. IT IS LOCATED BETWEEN IGG1 AND IGG2.
- 48) HUMAN IGG2'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN GENOMIC DNA.
- 53) HUMAN IGG4'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN GENOMIC DNA.
- 55) HUMAN IGC'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCES OF CLONES OF HUMAN GENOMIC DNA FROM FETAL LIVER (NISHIDA, Y., MIKI, T., HISAJIMA, H. & HONJO, T. (1982) PROC. NAT. ACAD. SCI. USA, 79, 3833-3837) AND PLACENTA (MAX, E.E., BATTLEY, J., NEY, R., KIRSCH, I.R. & LEDER, P. (1982) CELL, 29, 691-699), AND FROM A HUMAN IGE PRODUCING MYELOMA 6681 (FLANAGAN, J.S., RABBITTS, S. H. (1982) EMBO J., 1, 655-660). ALL THREE SEQUENCES HAVE LEU AT POSITION 539. IN ADDITION, AT POSITION 273 NOT REPORTED BY AMINO ACID SEQUENCING IN IGE MYELOMA NO. MAX ET AL. AND FLANAGAN & RABBITTS CARRIED OUT COMPLETE NUCLEOTIDE SEQUENCES AND NISHIDA ET AL. PARTIAL SEQUENCES EXCEPT FOR RESIDUE 539 WHICH IS REPORTED AS LEU BY MAX ET AL. AND AS TRP BY FLANAGAN & RABBITTS. ALL AMINO ACID RESIDUES ARE IDENTICAL.
- 56) HUMAN IGC'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF HUMAN CDNA.
- 56) IGM'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF NEWBORN MOUSE DNA. THE LENGTHS OF INTERVENING SEQUENCES BETWEEN CH1 AND HINGE, HINGE AND CH2, AND CH2 AND CH3 ARE 110, 279, AND 107 BASE-PAIRS RESPECTIVELY. HONJO, T., MATSUZAKI, S., RABBITTS, S. H. (1980) NUCLEIC ACIDS RES. 8, 703-713. HAVE ALSO OBTAINED AMINO ACID RESIDUES 583 TO 628 FROM THE NUCLEOTIDE SEQUENCES. CALAME, K., ROGERS, J., EARLY, P., DAVIS, M., LIVANT, D., WALLER, R., HODGE, J. (1980) NATURE, 284, 425-428. BY NUCLEOTIDE SEQUENCING OF AMINO ACIDS 320 TO 343 AND 593 TO 628 FOUND ACTUALLY FOR CH1 AT POSITIONS 111-125.
- 67) IGM'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA. A SINGLE AMINO ACID SUBSTITUTION AT POSITION 435 FROM SER TO ASN RESULTS IN A MUTANT PROTEIN WHICH IS DEFECTIVE IN INITIATING COMPLEMENT-DEPENDENT CYTOLYSIS. (SHULMAN, M.J., PENNELL, N., COLLINS, C. & HAZUM, N. (1986) PROC. NAT. ACAD. SCI. USA, 83, 7678-7682).
- 69) MUTANT 102'CL: IT LACKS CODONS 601 TO 614, SO THAT PREDOMINANTLY MONOMERIC Igm WAS PRODUCED BY THIS CELL LINE.
- 71) MOPC104E MOXB'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CDNA CLONE.
- 72) EPC76'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CDNA CLONE.
- 75) IGD'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE DNA.
- 76) IGD SECR'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE GENOMIC DNA.
- 77) IGD MOXB'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE GENOMIC DNA.
- 81) IGG1'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF NEWBORN MOUSE DNA. THE LENGTHS OF INTERVENING SEQUENCES BETWEEN CH1 AND HINGE, HINGE AND CH2, AND CH2 AND CH3 ARE 110, 279, AND 107 BASE-PAIRS RESPECTIVELY. HONJO, T. & RABBITTS, S. H. (1980) NUCLEIC ACIDS RES. 8, 703-713. HAVE DETERMINED RESIDUES 114 TO 128, 313 TO 328, AND 335 TO 376. THEY ARE IDENTICAL AT CORRESPONDING POSITIONS.
- 82) IGG1 MOXB'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA.
- 84) I72'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA FROM CELL LINE I72. THE CHI REGION IS DELETED.
- 86) IGG2B(A)'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA. THE LENGTHS OF INTERVENING SEQUENCES BETWEEN CH1 AND HINGE, HINGE AND CH2, AND CH2 AND CH3 ARE 316, 108, AND 112 BASE-PAIRS RESPECTIVELY.
- 87) IGG2B(A)'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA. THE LENGTHS OF INTERVENING SEQUENCES BETWEEN CH1 AND HINGE, HINGE AND CH2, AND CH2 AND CH3 ARE 316, 107, AND 112 BASE-PAIRS RESPECTIVELY.
- 88) IGG2B(B)'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA. THE LENGTHS OF INTERVENING SEQUENCES BETWEEN CH1 AND HINGE, HINGE AND CH2, AND CH2 AND CH3 ARE 317, 108, AND 113 BASE-PAIRS RESPECTIVELY.
- 89) IGG2B MOXB'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA.
- 90) MOPC11: THE AMINO ACID RESIDUES AT POSITION 363 WERE FOUND TO BE ILE AND LEU.
- 10.1: THIS PROTEIN IS OBTAINED FROM A MUTANT OF MOPC11 WITH A DELETION OF 99 NUCLEOTIDES INCLUDING THE 3' END OF THE CHI EXON, GIVING RISE TO A DELETION OF THE ENTIRE CHI REGION OF THE PROTEIN.
- 92) IGG2A(A)'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA.
- 93) I7/9'CL: THE Fc2 FRAGMENT OF I7/9 (IGG2a-KAPPA) HAS BEEN CRYSTALLIZED.
- 94) IGG2A(B)'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA.
- 95) IGG2A MOXB'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA.
- 96) IGA'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE CDNA.
- 97) IGA'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF BALB/C MOUSE LIVER DNA. THE HINGE REGION OF THE HINGE REGION IS NOT ON A SEPARATE EXON. THE FIRST THREE AMINO ACID RESIDUES BELONG TO THE CHI EXON, AND THE REMAINING NINE AMINO ACID RESIDUES BELONG TO THE CH2 EXON. THEY HAVE BEEN PLACED IN THE HINGE DOMAIN SINCE THEY ALIGN PERFECTLY WITH OTHER IGAS.
- 100) MOPC511: CARBOHYDRATES ARE ATTACHED AT POSITIONS 154 AND 483.
- 101) IGC'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF MOUSE DNA FROM AN IGE-PRODUCING HYBRIDOMA.
- 102) IGE a'CL: FROM BALB/C MOUSE LIVER DNA.
- 103) IGE b'CL: FROM BALB/C MOUSE LIVER DNA.
- 104) IGE MOXB'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF DNA FROM NEWBORN MOUSE.
- 105) RAT IGD'CL: THE AMINO ACID SEQUENCE WAS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF RAT CDNA.
- 106) I72'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF RAT CDNA.
- 107) IR-162'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF RAT CDNA.
- 108) PGAMGAB1-12.14'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF RABBIT CDNA.
- 109) RAB1GG: THE AMINO ACID SEQUENCE FOR POSITIONS 266 TO 275 AND 321 TO 336 WAS ALSO CONFIRMED BY OTHERS (TEHERANI, J., CAPRANICO, AGGARWAL, S. & MANDY, W.J. (1979) EUR. J. IMMUNOL. 9, 690-695).
- 110) P2A2'CL: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE NUCLEOTIDE SEQUENCE OF A CLONE OF RABBIT CDNA.

## SPECIFIC NOTES: HEAVY CONS

- 143) SEKED pSBC'CL: TRANSLATED.
- 144) BOREE IGD: AS COMPARED WI
- 147) CHICKEN IGM'CL: THE AMINO
- 148) Slope VM'CL: FROM Elops sa
- 152) *Xenopus laevis* c8(II)'CL:
- 153) *Xenopus laevis* c14(II)'CL:

- THE FOLLOWING WERE EQUALLY A

AT POSITION		(ILE
113D		
113E		
113F		
113G		
224		
241B		
231I		
213		
307		
326		
313		
351A		
403		
482		
491C		
551		
512		
600		
606B		
617		
615		
606		
606		

**SPECIFIC NOTES: HEAVY CONSTANT CHAINS** (cont'd)

- 143) **SHEEP pSHC'CL**: TRANSLATED FROM cDNA OF SHEEP LYMPHOCYTES  
 144) **HORSE IGG**: AS COMPARED WITH HORSE IGG, THE HORSE T PROTEIN HAD VAL AT POSITION 463, GLU AT 464, AND HIS AT 474.  
 147) **CHICKEN IGM'CL**: THE AMINO ACID SEQUENCE IS OBTAINED BY TRANSLATING THE SEQUENCE OF A CLONE OF CHICKEN CDNA.  
 148) **Elops VH'CL**: FROM Elops saurus (LADYFISH).  
 152) **Xenopus laevis c8(II)'CL**: ALSO KNOWN AS XIg8'CL.  
 153) **Xenopus laevis c14(II)'CL**: ALSO KNOWN AS XIg14'CL.

+ THE FOLLOWING WERE EQUALLY AND MOST FREQUENTLY OCCURRING:

AT POSITION	RESIDUES
113D	(ILE, GLN), (ILE, GLU)
113E	(CYS, SER)
137	(THR, SER)
158	(LYS, ASN)
224	(ARG, GLU)
241B	(PRO, CYS)
243I	(ALA, SER)
281	(PRO, GLU)
307	(PRO, LYS)
326	(PRO, THR)
343	(THR, ASN)
351A	(LEU, THR, ASN)
401	(THR, ASN)
452	(ASP, ASN)
496C	(PRO, ALA)
511	(ARG, GLU)
532	(ALA, GLU)
545	(PRO, LEU)
580B	(MET, ASP)
637	(LEU, VAL)
657	(GLU, GLN)
662	(ILE, VAL, ALA)
663	(ILE, SER)

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